

[1,2,4]-TRIAZOLO[1,5-c]PYRIMIDINE DERIVATIVE**Publication number:** WO2004029056**Publication date:** 2004-04-08**Inventor:** IIDA KYOICHIRO (JP); SUGITA TAKAMASA (JP); SHIOZAKI SHIZUO (JP); KANDA TOMOYUKI (JP); KUWANA YOSHIHISA (JP); SHIMADA JUNICHI (JP)**Applicant:** KYOWA HAKKO KOGYO KK (JP); IIDA KYOICHIRO (JP); SUGITA TAKAMASA (JP); SHIOZAKI SHIZUO (JP); KANDA TOMOYUKI (JP); KUWANA YOSHIHISA (JP); SHIMADA JUNICHI (JP)**Classification:**

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Cited documents:

- WO9503806
- EP0459702
- EP0515107
- WO9842711
- WO0017201
- WO03068776
- XP002974911

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A [1,2,4]-triazolo[1,5-c]pyrimidine derivative represented by the general formula (I): (I) (wherein R<1> represents (un)substituted aryl or an (un)substituted aromatic heterocyclic group; R<2> represents hydrogen, halogeno, lower alkyl, lower alkanoyl, aroyl, (un)substituted aryl, or an (un)substituted aromatic heterocyclic group; R<3> represents lower alkyl, lower cycloalkyl, (un)substituted lower alkanoyl, (un)substituted aryl, an (un)substituted aromatic heterocyclic group, etc.; and Q represents hydrogen or 3,4-dimethoxybenzyl) or a pharmacologically acceptable salt of the derivative. They have antagonistic activity on an adenosine A2A receptor and are useful for treatments for and/or prevention of diseases attributable to adenosine A2A receptor hyperenergia.

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